

**K936310 ORTHOMET HMC FEMORAL STEM COMPONENT**Jun 7, 1995  
426 days to decisionK936310 · Product code: **LPH** · Orthopedic  
Source: <https://www.510kdatabase.net/k936310/>**SUBMISSION DETAILS**

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|                       |   |
|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)  |
| Submission type       | Traditional   |
| Device classification | Prosthesis, Hip, Semi-constrained, Metal/polymer, Porous Uncemented (LPH) |
| Date received         | Apr 7, 1994   |
| Decision date         | Jun 7, 1995   |
| Days to decision      | 426 days  |
| Third-party review    | No  |
| Summary / Statement   | Summary   |

**APPLICANT**

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|----------------|---|
| Company        | <b>Orthomet, Inc.</b>                   |
| Location       | Plymouth, MN, US                        |
| Contact        | DAVID CANNISTRACI                       |
| 510(k) history | 60 submissions · 41 cleared · 1986-1995 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k936310/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 21, 2026